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Title

Lagenidium giganteum pathogenicity in mammals.

Permalink

<https://escholarship.org/uc/item/3z68443z>

Journal

Emerging infectious diseases, 21(2)

ISSN

1080-6040

Authors

Vilela, Raquel
Taylor, John W
Walker, Edward D
et al.

Publication Date

2015-02-01

DOI

10.3201/eid2102.141091

Peer reviewed

Lagenidium giganteum Pathogenicity in Mammals

Raquel Vilela, John W. Taylor,¹ Edward D. Walker,¹ Leonel Mendoza

Infections of mammals by species in the phylum Oomycota taxonomically and molecularly similar to known *Lagenidium giganteum* strains have increased. During 2013–2014, we conducted a phylogenetic study of 21 mammalian *Lagenidium* isolates; we found that 11 cannot be differentiated from *L. giganteum* strains that the US Environmental Protection Agency approved for biological control of mosquitoes; these strains were later unregistered and are no longer available. *L. giganteum* strains pathogenic to mammals formed a strongly supported clade with the biological control isolates, and both types experimentally infected mosquito larvae. However, the strains from mammals grew well at 25°C and 37°C, whereas the biological control strains developed normally at 25°C but poorly at higher temperatures. The emergence of heat-tolerant strains of *L. giganteum* pathogenic to lower animals and humans is of environmental and public health concern.

During the 20th century, *Pythium insidiosum* was the only fungus-like species from the phylum Oomycota known to cause life-threatening infections in mammals and birds (1,2). In 2003, a novel group of pathogenic Oomycota was isolated in the southeastern United States from dogs with invasive cutaneous infections resembling pythiosis, but the strains had morphologic and molecular affinities with the genus *Lagenidium* (3,4). Subsequently, other strains similar to previously studied *Lagenidium* species were found to be associated with infections in dogs, cats, and humans (5–7). The disease caused by these 2 Oomycota species has been reported mainly in apparently healthy hosts; thus, these species' evolutionary traits need to be investigated (1–7). Although detailed descriptions of the clinical, pathologic, and serologic features of *Lagenidium* spp. pathogenic to mammals have been published (3,4), their taxonomic relationship to *Lagenidium* strains approved for biological control and the capacity of their zoospores to infect mosquitoes have yet to be investigated (5,6).

The genus *Lagenidium*, introduced by Schenk in 1857 (8), comprises numerous saprotrophic species (9), as well as

several members that are pathogenic to algae, phytoplankton, pollen, barnacles, blue crabs, mosquito larvae, shrimp, and mammals (3–5). Couch (10) isolated *L. giganteum* from *Daphnia* larvae in Virginia, USA. Later, Couch and Romney (11) confirmed that *L. giganteum* infects and kills mosquito larvae as it reproduces sexually. Their work stimulated interest in *L. giganteum* as a biological control agent of mosquitoes (11,12). In 1995, the US Environmental Protection Agency registered *L. giganteum* under the trade name Lagenex as a biocontrol agent (13) but later deregistered it at the request of the manufacturer (<http://www.gpo.gov/fdsys/pkg/FR-2011-09-28/pdf/2011-24832.pdf>). *L. giganteum* strains used for testing and commercial development were deposited within the American Type Culture Collection (ATCC), and the ATCC strains and other not-well-characterized strains were released in nature for field studies (12–16). In this study, conducted during 2013–2014, we investigated 21 mammalian *Lagenidium* strains and compared them with *L. giganteum* isolates used for mosquito control.

Materials and Methods

Living Cultures

A complete list of the strains and their accession numbers is shown in the Table (<http://wwwnc.cdc.gov/EID/article/21/2/14-1091-T1.htm>). In brief, we studied the following isolates: *L. giganteum* (ATCC 36492 = ATCC 52675 used as biological control and ATCC 48336 isolated from mosquito larvae); *L. humanum* (ATCC 76726 isolated from the skin of dead humans); *L. giganteum* pathogenic to mammals (culture collection at the Biomedical Laboratory Diagnostics, Michigan State University (MTLA): MTLA 01 = ATCCMYA-4933 type strain, MTLA-03, MTLA-04 = ATCCMYA-4934 (from a US cat), MTLA-05 = ATCCMYA-4935, MTLA-10, MTLA-12, MTLA-14, MTLA-15, MTLA-16, MTLA-17, and MTLA-18 from dogs); *L. ajelloi* (MTLA-06 = ATCCMYA-4936 type strain, MTLA-07 = ATCCMYA-4937, MTLA-19, MTLA-20, MTLA-21, MTLA-22, MTLA-23, isolated from dogs); *L. vilelae* (MTLA-24 = DNA extracted from fixed feline tissue and MTLA-25 cat strain); and *L. albertoi* (MTLA13 = ATCCMYA-4932 type strain).

Author affiliations: Michigan State University, East Lansing, Michigan, USA (R. Vilela, E.D. Walker, L. Mendoza); Universidade Federal do Minas Gerais, Belo Horizonte, Brazil (R. Vilela); University of California, Berkeley, California, USA (J.W. Taylor)

DOI: <http://dx.doi.org/10.3201/eid2102.141091>

¹These authors contributed equally to this article.

Media and Culture Conditions

The isolates were grown on brain-heart infusion (BHI) (DIFCO, Detroit, MI, USA), corn meal agar (CMA) (BBL, Sparks, MD, USA), 2% Sabouraud dextrose agar (SDA), 2% Sabouraud dextrose broth, in triplicate. Cardinal temperatures of growth (25°C, 30°C, and 37°C measured during 3 days' incubation) and their relation to the production of sexual and asexual structures were evaluated on the above media. Development of the different stages of vesicle and zoospore formation, cleavage, and release were assessed on colonized grass leaves in water cultures containing Ca^{++} and Mg^{++} . Briefly, the evaluated strains (Table) were subcultured on SDA plates at 37°C for 24 h. After incubation, 5 × 5-mm diameter blocks were cut from the advancing edges of the culture and placed on top of a 2% water agar plates. Sterile 4 × 10-mm grass blades were laid on top of each block and incubated at 37°C for 24 h (*L. giganteum* = MTLA-01 from mammals). The grass blades were then collected and placed in a beaker that contained 50 mL of sporulation solution (made of 2 solutions: mix no. 1 contained $\text{NH}_4\text{H}_2\text{PO}_4$ [66.04 g], KH_2PO_4 [68.05 g], and K_2HPO_4 [87.09 g] in 500 mL H_2O ; and mix no. 2 contained $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$ [18.38 g] and $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$ [25.42 g] in 250 mL H_2O). The sporulation solution was obtained by mixing 0.5 mL of solution no. 1 plus 0.1 mL of solution no. 2 in 1.0 L of distilled water. The beaker with the 50-mL of sporulation solution plus the parasitized grass blades was incubated at 37°C, and the development of sporangia and zoospores microscopically was evaluated every 30 min for the following 6 h (or longer when needed).

Morphologic Description of *L. giganteum* from Mammals

We evaluated the morphologic features of *L. giganteum* strains after subculture on BHI, CMA, and SDA at different intervals during 5 days' incubation at 37°C or at room temperature (25°C). Briefly, 4 × 4-mm agar blocks were cut from the above media and mixed with 1 drop of lactophenol cotton blue (phenol 20 mL, lactic acid 20 mL, glycerol 40 mL, and distilled water 20 mL). The morphologic features of their hyphal and other structures developed in these media, including oogonia, were microscopically investigated. We evaluated the development of sporangia and zoospores in sporulation medium (see above). After incubation at the induction temperatures, the beaker containing 50 mL of sporulation medium plus the parasitized grass blades was inspected on an inverted microscope. If vesicles and zoospores developed, we removed the grass blade with ≥10 vesicles, using tweezers, and placed it on a glass slide. Immediately, 5 µL of Merthiolate (0.02%) was added to the grass blade to stop the movement of the zoospores and facilitate measurements. Alternatively, grass blades containing vesicles with zoospores were also transferred to plates holding *Culex pipiens* larvae.

Experimental Mosquito Infection

The strains of *L. giganteum* from mammals (MTLA01, MTLA03, MTLA04, MTLA05, and MTLA10) (Table) were used to evaluate their capability to infect mosquito-to larvae of *C. pipiens*. Instar 3 *C. pipiens* larvae were collected and placed in 6-well plates (1 plate per experiment; Corning Inc., Corning, NY, USA) containing 5 mL of water and fed every other day with fish food flakes (TetraMin, Lancaster, PA, USA) until the end of the experiment. We transferred fresh grass blades, with ≥10 vesicles per blade containing numerous zoospores, to the plates holding 5 mosquito larvae per well (total of 30 larvae per plate). The plates were incubated at 25°C and observed daily by using an inverted microscope for the next 3 weeks. The criteria to select putative infected mosquito larvae were based on their swimming capabilities. We closely inspected slower swimmers for filamentous structures outside and between the larval segments and inside their bodies. Every trial included its positive (*L. giganteum* biological control ATCC 36492 = ATCC 52675) and negative controls (larvae without exposure to *L. giganteum* species), and the experiment was repeated twice (total 60 *C. pipiens* larvae).

DNA Extraction and Molecular Procedures

The 20 strains investigated (Table) were inoculated into 250-mL flasks containing 100 mL of Sabouraud dextrose broth and incubated for 72 h at 37°C on a shaker rotating at 150 rpm. In addition, 1 formalin-fixed tissue sample from a cat was also used to extract total genomic DNA following the company protocols (QIAGEN, Valencia, CA, USA). After incubation, the cultures were killed with Merthiolate (0.02%, wt/vol) and then filtrated to separate the hyphal cell mass. The hyphal cell mass was then transferred to a mortar and ground in the presence of liquid nitrogen. DNA from the disrupted hyphae was treated with sodium dodecyl sulfate and proteinase K and then incubated at 60°C for 1 h, and genomic DNA of the hydrae was extracted with phenol, chloroform, isoamyl alcohol (Sigma, St. Louis, MO, USA). PCR was conducted by hot start amplification using the following primers: the universal primers for internal transcribed spacers (ITS): ITS1 = TCCGTAGGTGAACCTGCGG and ITS4 = TCCTCC-GCTTATTGATATGC; cytochrome oxidase II (COXII): COX.LagF = 5'-CCACAAATTTCACCTACATTGA-3' and COX.LagR = 5'-TAGGATTTCAAGATCCT-GC-3'; heat shock protein 90 (HSP90): HSP90.LagF = 5'-CAACCTBGGHACSATYGCCAAG-3' and HSP90.LagR = 5'-ACRAAMGACARGTAYTCVGGCA-3'; cell division cycle 42 (CDC42): CDC42.LagF = 5'-GTSC-CVACYGTVTTYGANAAAYTA-3' and CDC42.LagR = 5'-GCWSWGCAITCVASRTAYTT-3'; Tubulin: TUB.LagF = 5'-GGTGGTGGTACCGGTTTC-3' and TUB.LagR

= 5'-GACACACGCTTGAACATC-3'. In addition, the following primers were constructed to PCR amplify some of the *L. ajelloi* and *L. vilelae* strains: CDC42.Lag2F = 5'-GTGCCGACYGTGTTYGABAAC-3' and CDC42.Lag2R = 5'-CTSTTCKGTTGTRATBGG-3'; and HSP90.Lag2F = 5'-GAGGCCTTCGTGGAAGCG-3' and HSP90.Lag2R = 5'-GTTGTTTCATTTTCTTGCGGG-3'. The PCR temperature-cycling parameters were as follows: 10 min

at 95°C and 1 min for subsequent cycles, annealing for 1 min at 60°C, and elongation at 72°C for 2 min. The parameter was repeated for 40 cycles, followed by a final elongation of 2 min at 72°C for 7 min. The amplicons were ligated into pCR 2.1-TOPO vector (Invitrogen, Carlsbad, CA, USA), purified, and then sequenced by using BigDye Terminator chemistry in an ABI Prim 310 genetic analyzer (Perkin-Elmer, Foster City, CA, USA).

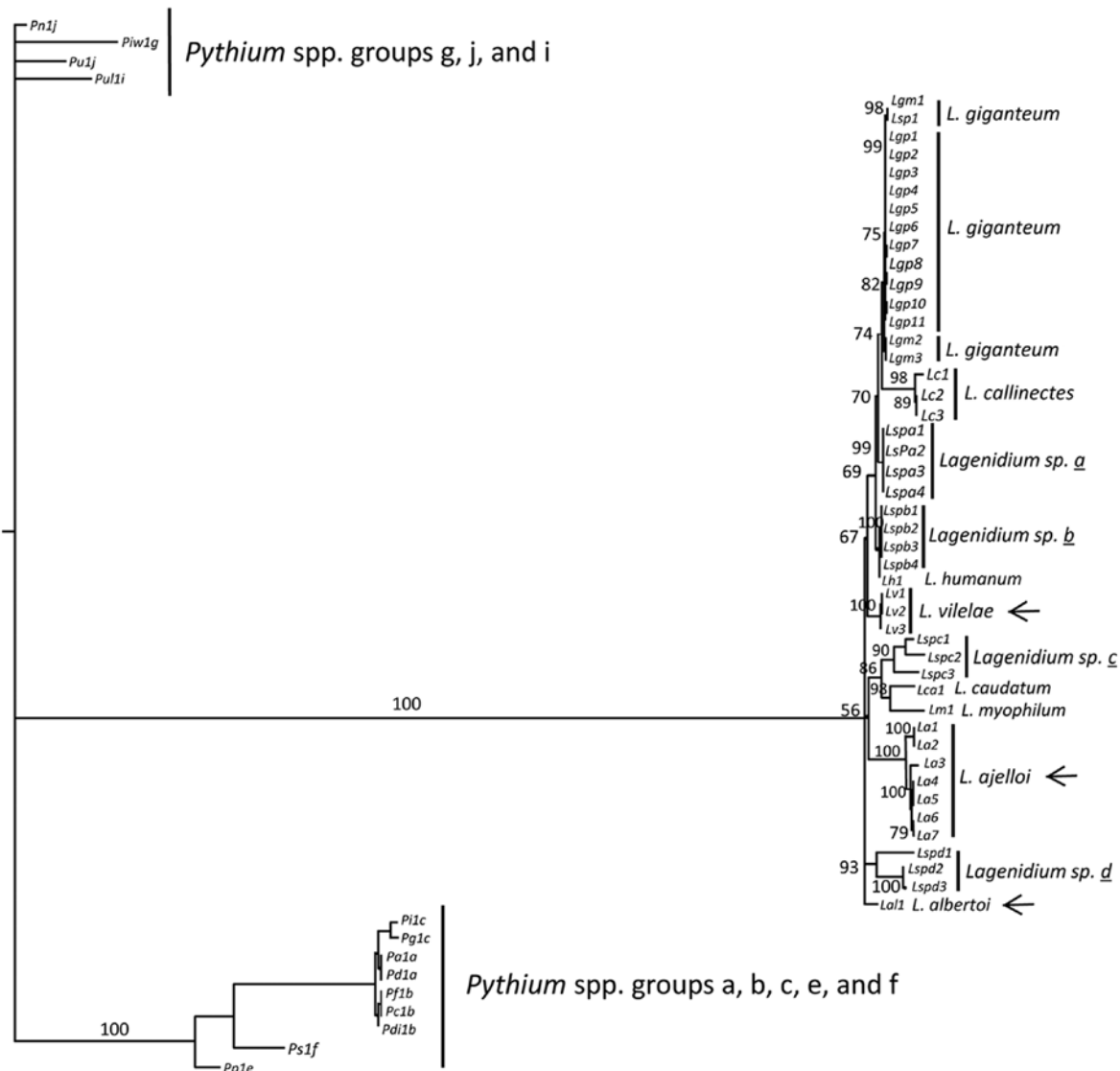


Figure 1. Bayesian phylogenetic analysis of concatenated 4 partial coding gene sequences (cell division cycle 42, cytochrome oxidase II, heat shock protein 90, and tubulin) and the complete internal transcribed spacers 1, 2, and 5.8S of *Lagenidium* DNA sequences. Thirteen *Pythium* species DNA sequences were included as the outgroup (groups a–c, e–g, j, and i [16]; Table, <http://wwwnc.cdc.gov/EID/article/21/2/14-1091-T1.htm>). Support on key branches is the Bayesian probability for that branch followed by the percentage of 1,000 bootstrap resampled datasets containing the branch in neighbor-joining analyses of maximum-likelihood distances followed by the percentage of 1,000 bootstrap resampled datasets containing the branch in parsimony analyses using heuristic searches. In this analysis, the DNA sequences of *L. giganteum* mosquito control (Lg 1–3) and a *Lagenidium* sp. recovered from a nematode in Taiwan (Ls1, Lsp1 = HQ395647) clustered with *L. giganteum* from mammals (Lg 1–10). The pathogen of crab *L. callinectes* (Lc) was the sister group to the cluster. Three *Lagenidium* mammalian pathogenic novel species (*L. ajelloi* = La, *L. albertoi* = Lal, and *L. vilelae* = Lv) were placed in 3 distinctive strongly supported clades (arrows). The accession numbers, the abbreviations used to identify each species, and the *Lagenidium* and *Pythium* spp. DNA sequences are shown in the Table. ATCC, American Type Culture Collection; CBS, Centraalbureau voor Schimmelcultures.

Phylogenetic Analysis

Genomic DNA sequences of the studied strains' complete ITS and the partial gene sequences of the selected 4 exons (*CDC42*, *COXII*, *HSP90*, and *TUB*) were aligned with other *Lagenidium* and *Pythium* DNA sequences available at the National Center for Biotechnology Information by using ClustalW version 1.81 (<http://www.clustal.org>) with default settings and their alignments visually inspected. *Pythium* spp. groups a–c, e–g, and j (15) (Table) were used as outgroups. Phylogenetic and molecular evolutionary analyses were conducted by using MEGA6 (<http://www.megasoftware.net>). The aligned sequences were exported for parsimony analysis by using a heuristic search with tree bisection reconnection branch swapping (MEGA6) and distant analysis by neighbor-joining (MEGA6). We coded

large insertions as 1 event by excluding all but 1 nt per insertion. The generated gaps were treated as missing data. Neighbor-joining analysis used either uncorrected distances or maximum-likelihood estimates of distance with a time-reversible model (6ST), and empirical base frequencies with no rate variation among sites, or a shape parameter of 0.5 γ -distribution with 4 rate categories. Branch support was estimated as percentage of parsimony tree (1,000 resampling, heuristic, branch swapping) or neighbor-joining trees (1,000 resampling, maximum-likelihood distances) within each branch. Concatenated DNA sequences of the 5 loci investigated were used in Bayesian analysis. We conducted the test in MrBayes version 3.2.1 $\times 64$ (<http://mr bayes.sourceforge.net>) using the general time reversible + I + γ model, with 2 chains (1 heated), 2 runs, sampling

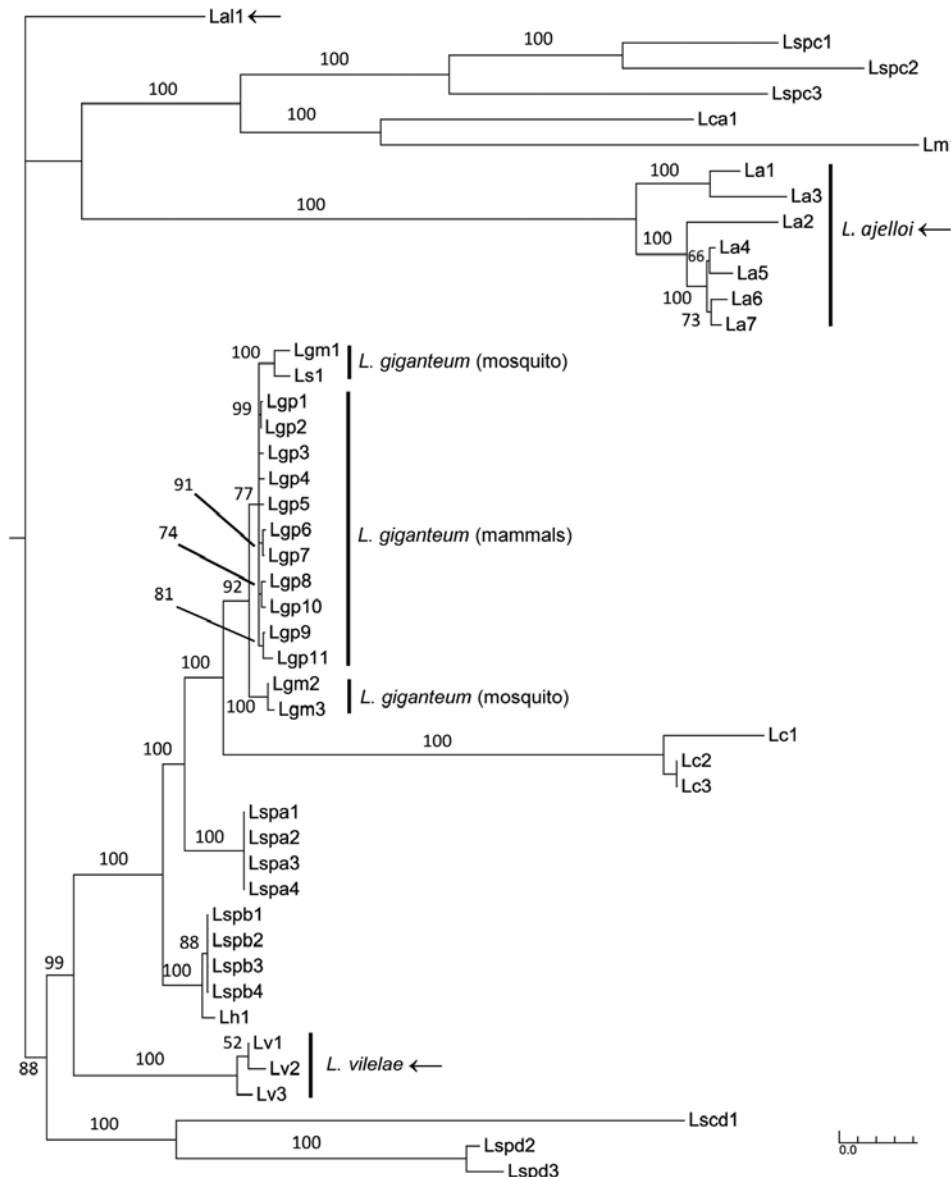


Figure 2. The Bayesian tree was constructed by concatenated aligned of *Lagenidium* spp. DNA sequences as in Figure 1 without outgroup to highlight the position of *L. giganteum* in the tree. *L. giganteum* from mammals (Lgp 1–11), *L. giganteum* mosquito control (Lgm 1–3 and Ls1 = HQ395647), and the novel species *L. ajelloi* = La, *L. albertoi* = Lal, and *L. vilelae* = Lv were placed in 4 strongly supported clades (arrows). Scale bar indicates nucleotide substitutions per site.

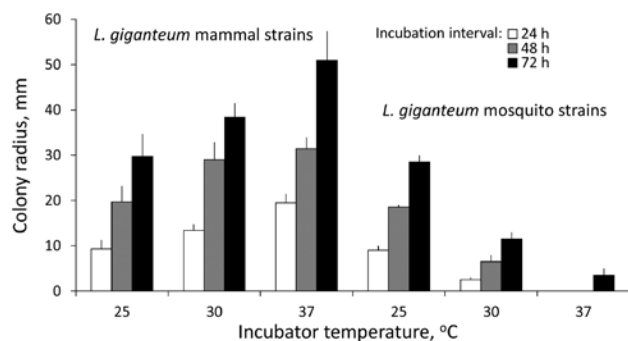


Figure 3. Cardinal temperatures of *Lagenidium giganteum* types in culture. Growth (mean colony radius and SEM, mm) of *L. giganteum* mammalian and mosquito strains at 3 temperatures at 24-, 48-, and 72-hour intervals postinoculation onto 2% Sabouraud dextrose agar. Repeated measures analysis of variance showed highly significant differences between strains ($F_{1,33} = 165.0$, $p < 0.0001$) and a highly significant interaction of strain and incubation temperature across time intervals ($F_{2,33} = 45.9$, $p < 0.0001$).

every 100th generation for 1×10^6 generations, and exclusion of the first 2.5×10^5 samples (the burn-in) before analysis. Support for branches was estimated as the percentage of parsimony tree (1,000 resampling, heuristic, nni branch swapping) or neighbor-joining tree (1,000 resampling, maximum-likelihood distances) containing the branch, as well as by determining the Bayesian probability estimated as the percentage of Bayesian trees possessing the branch after discarding the burn-in sample.

Results

Phylogenetic Analysis

Using *Pythium* spp. groups a–c, e–g, and j (Table) as outgroups, we sorted the concatenated DNA sequences of 4 protein-coding genes and 1 ribosomal region in Bayesian and maximum-likelihood phylogenetic analyses 21 mammalian pathogenic *Legenidium* strains into 4 strongly supported clades in the combined analysis (Figure 1). In the largest of these clades, 11 of the 21 strains (Table) recovered from infected mammals form a monophyletic clade that includes other *L. giganteum* strains previously used to develop biological control agents of mosquitoes (Figure 1). Also included in this clade is an *L. giganteum* strain recovered from nematodes in Taiwan (Lsp1). The remaining 10 mammal-infecting *Legenidium* strains were resolved with strong support into 3 independent clades that are basal to *L. giganteum* and identified in this study as *L. ajelloi*, *L. albertoi*, and *L. vilelae* (Figures 1, 2). These new species will be fully described elsewhere. Other clades of *Legenidium* species, as yet unnamed, are also apparent in the phylogeny (Lspa to Ld; Figures 1, 2).

Heat-tolerance and Mosquito Larva

Infection Experiments

The 11 *L. giganteum* strains from mammalian hosts grew faster at 37°C and 30°C than at 25°C, whereas *L. giganteum* biological control strains (ATCC 36492 = ATCC 52675 and ATCC 48336) grew well at 25°C and poorly at 30°C and almost failed to develop at 37°C in growth experiments conducted in CMA and SDA media (Figure 3). The biological control strains barely developed at 37°C but survived, as shown by their subsequent growth, after being transferred to 25°C.

We tested the ability of the mammalian *L. giganteum* zoospores to infect mosquito larvae in 5 *L. giganteum* mammal strains (MTLA01, 03, 04, 05, and 10; Table) and 2 *L. giganteum* biocontrol strains (ATCC 48336 = Lgm2, ATCC 36492 = Lgm3; Table). The emerging mammalian pathogenic strains exhibited mosquito-infection capabilities similar to those of the strains approved by the US Environmental Protection Agency for mosquito control (Figure 4). At least 1 larvae per well was found infected. Infected larvae exhibited abnormal swimming behavior and died within 2 days (Figure 4). Of the 60 *C. pipiens* larvae (2 trials, 30 larvae per experiment), 12 were found infected at the end of both experiments (instar 3 or instar 4). Uninfected larvae continued their normal life cycle. Postmortem examination of infected larvae showed extensive infections with numerous hyphae spreading internally and emerging between segments (Figure 4, panels A–E). Under our experimental conditions, *L. giganteum* ATCC 36492 infected 18 of the 60 *C. pipiens* larvae. In unexposed controls all larvae survived, larvae swam normally, and we found no evidence of infection. The identity of mammalian *L. giganteum* strains experimentally infecting mosquito larvae was confirmed by culture, DNA extraction, sequencing, and phylogenetic analysis at the conclusion of each experiment.

Morphologic Characteristics of *L. giganteum* in Culture

Submerged colonies of both *L. giganteum* types (biological control and mammalian strains) are colorless or yellow with few aerial mycelia that present irregular radiating and undulating patterns on any of 3 media (CMA, BHI, or 2% SDA) (Figure 5, panels A, D). Typically, the elongated mycelium comprises ovoides or spherical 30- to 40-μm diameter segments forming chains and connected by broad to slender isthmuses; hyphae also form long irregular segments of ≥ 400 μm (Figure 5, panels B, C, E, F). We did not find oogonia in the evaluated strains. In liquid sporulation medium, hyphal segments develop 1–2 discharge tubes that produce terminal vesicles containing reniform zoospores 9–10 μm wide \times 10–12 μm long (Figure 5, panel G). Zoospores were released in both mammal and mosquito *L. giganteum* strains after the discharge vesicle was disrupted (Figure 5, panel H), which then swam for 10–20 minutes

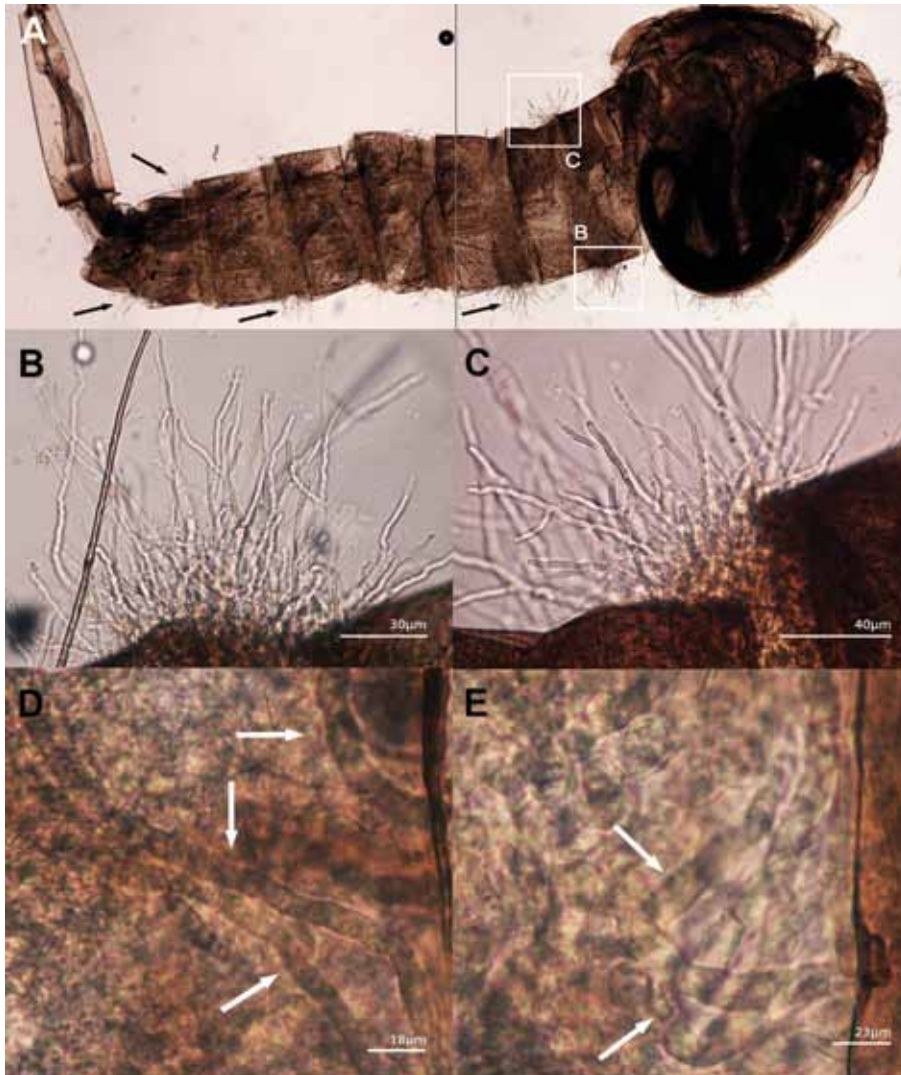


Figure 4. *Lagenidium giganteum* from mammal experimental infection using *Culex pipiens* mosquito larvae. A) Composite of 2 photographs showing an instar 3 *C. pipiens* larvae infected with 1 of the 5 tested strains of *L. giganteum* recovered from dogs with lagenidiosis (MTLA01, type strain). Note the mycelioid structures emerging from the infected larvae (arrows). B, C) Enlargements of the 2 white boxes in (A) showing details of the mycelioid structures emerging between the segments of the larvae. D, E) Aggressiveness of the invading mycelioid structures within the body of *C. pipiens* larvae.

before encysting. Germ tubes developed from encysted zoospores within 10–20 minutes after encystment.

Discussion

We detected a novel group of mammalian pathogenic *L. giganteum* strains unique for their capacity to experimentally infect invertebrate hosts and cause life-threatening infections in healthy lower animals and humans (5,7). Placement of both heat-sensitive (affecting mosquito larvae) and heat-tolerant (affecting mammals) *L. giganteum* strains in a strongly supported monophyletic clade in Bayesian analysis corroborates the taxonomic and phenotypic attributes shared by both types in this and other studies (3–5,10–15). In addition to *L. giganteum*, 3 other clades harbor mammal isolates: *L. ajelloi*, *L. albertoi*, and *L. vilelai* (Figures 1, 2). Strains in the remaining clades are from invertebrates, whether from nematodes (the 4 unnamed species, *L. spa*, *L. spb*, *L. spc*, and *L. spd* [Table]),

crab eggs (*L. callinectes*), or shrimp (*L. myophilum*) (17). *Lagenidium* strains in the 3 clades closest to *L. giganteum* infect invertebrates, as can all isolates of *L. giganteum*, suggesting that the mammal-infecting *L. giganteum* isolates recently evolved the ability to infect warm-blooded hosts. However, the capacity to infect mammals is not unique to *L. giganteum*, and knowing that each of the 3 other mammal-infecting *Lagenidium* clades has closely related relatives that infect invertebrates, it seems likely that the trait of infecting mammals has arisen several times, independently, in the genus *Lagenidium*. From its name, 1 species, *L. humanum*, might be expected to be among the mammal parasites, but it has been isolated only from the dead skin of humans or snakes and cannot grow at 37°C (9). Regarding the trait of heat tolerance, species that are isolated from mammals, *L. ajelloi*, *L. albertoi*, *L. giganteum*, and *L. vilelai* also can grow at 37°C, but judging from cultures recovered from lower animals,

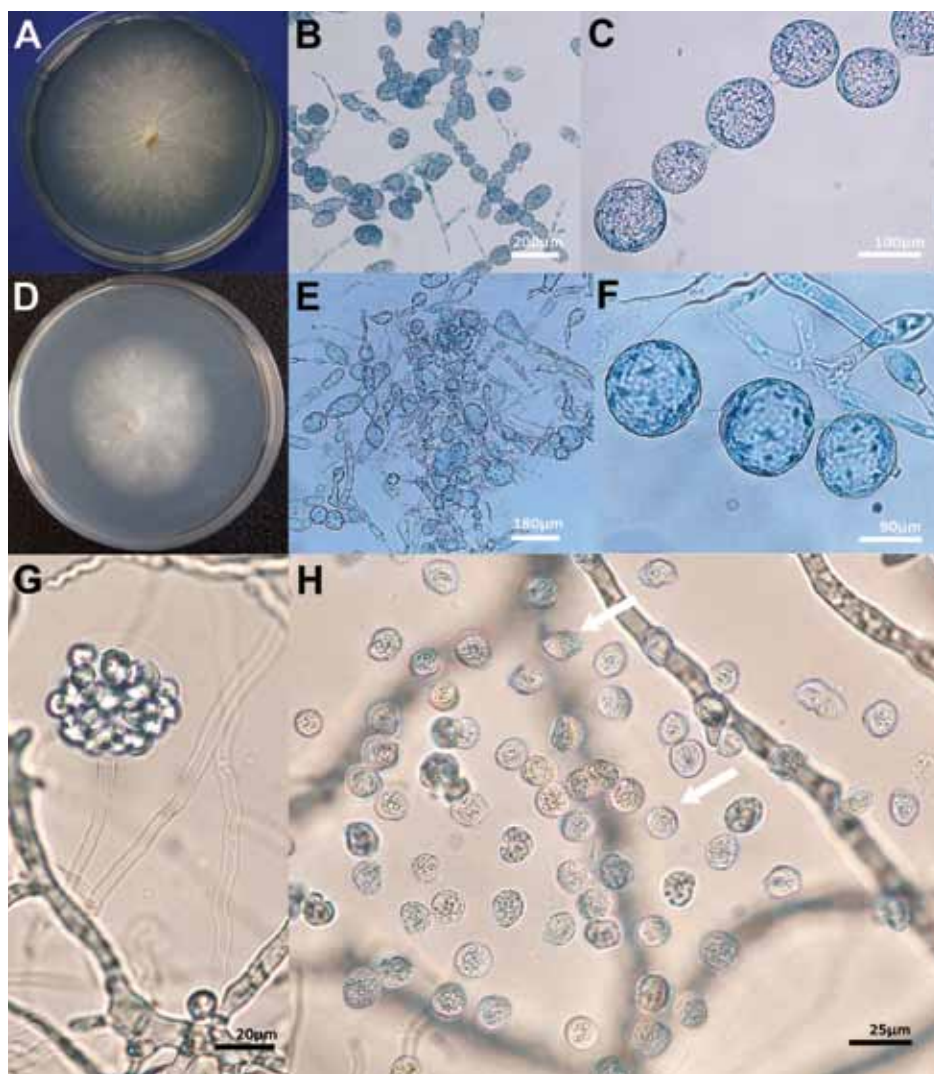


Figure 5. Morphologic features of isolates of *Lagenidium giganteum* mosquito control agent and *L. giganteum* mold from mammals. Panel A shows phenotypic features in culture of the mammalian pathogen (ATCCMYA-4933, type strain), and panel D shows the biological control (ATCC 36492). The development of spherical and ovoid 40- to 170-µm swelling segments (panels B, C, E, F) was the main feature of both mammalian and biocontrol strains. Panel G shows a tubular body developed from an unseen segmented sporangium form a vesicle enclosing numerous zoospores in a mammalian *L. giganteum* strain. The kidney-shaped zoospores before release (G) and after release (H, arrows) agree with those in the original description of *L. giganteum* by Couch (10). ATCC, American Type Culture Collection.

some strains associated with invertebrates also might lack the heat-tolerance trait. *Lagenidium* strain Centraalbureau voor Schimmelcultures (CBS) 127042 recovered from nematodes and phylogenetically linked to *L. vilelae* may be a good example of this unusual feature (Figure 1).

Emergence of mammalian pathogens often is accompanied by host switching in a zoonotic context from 1 vertebrate species to another (18,19), but range extension from invertebrates to vertebrates as hypothesized here is rare. Waterfield et al. (20) outlined 3 scenarios in which bacteria pathogenic to humans might have evolved from pathogens associated with invertebrates, but the time for the process was thousands of years, whereas in our second scenario, we describe a process that might have happened within a decade. The 2 phenotypes of heat-tolerance and mammal pathogenicity appear to have repeatedly evolved in the genus *Lagenidium*, and several recent studies have shown that members of Oomycota can acquire pathogenicity genes by

horizontal gene transfer (21–23). Before their release as agents of biocontrol, *L. giganteum* (ATCC 52675 = ATCC 36492) (12) and other strains proposed as biocontrol agents of insect were shown not to infect other arthropods and mammals (14,15). However, experimental infection of mammals by the other Oomycota pathogen, *P. insidiosum*, is difficult to achieve, e.g., only rabbits have been successfully, experimentally infected by this pathogen of cats, cattle, dogs, horses, and humans (2). This experience indicates that the safety of *Lagenidium* species as pathogens of hosts other than mosquito might be difficult to assess because of a general difficulty over experimental infection of mammals by Oomycota species.

The evolutionary relationships among *L. giganteum* strains should be more thoroughly examined by using population genomics, which could lead to discovery of the basis of adaptation to specific hosts, as has been shown for other adaptive phenotypes in other filamentous microbial eukaryotes (24).

The recent transcriptome analysis of a strain of *L. giganteum* recovered in nature from mosquito larvae further strengthens the relevance of our findings (25). The use of varieties or subspecies to name these 2 populations of *L. giganteum* was avoided until more comprehensive genomic data shed light on their true evolutionary relationships. Future genomic studies also could discriminate between the scenarios of gaining novel traits in specific ecologic niches or revealing types with hidden capabilities, both of which are of environmental and public health concern when the traits include putative pathogenicity of humans.

Acknowledgments

We thank the many veterinary practitioners and physicians who provided clinical samples from putative cases of lagenidiosis. We also thank R. Morris for comments on the manuscript.

This study was supported by the Department of Microbiology and Molecular Genetics, Michigan State University, and by grants R37AI21884, National Institute of Allergy and Infectious Diseases to E.D.W. and National Science Foundation DEB-125752 to J.W.T.

Dr. Vilela is a physician with a PhD in medical microbiology who works with unusual infectious fungal and fungal-like microbes at Biomedical Laboratory Diagnostics, Michigan State University. Her research interests include *Lacazia loboi*, *Lagenidium* spp., *Pythium insidiosum*, and *Rhinosporidium seeberi*.

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Address for correspondence: Leonel Mendoza, Biomedical Laboratory Diagnostics, 354 Farm Ln, Rm N322, Michigan State University, East Lansing, MI 48824-1031, USA; email: mendoza9@msu.edu

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